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APPLICATION NO. FILING DATE		FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/003,132 11/15/2001		Brian A. Fox	00-62	5882	
75	90 11/01/2002				
Gary E. Parker			EXAMINER		
ZymoGenetics, 1201 Eastlake A	venue East		NICHOLS, CHRISTOPHER J		
Seattle, WA 98102			ART UNIT	PAPER NUMBER	
			1647		
			DATE MAILED: 11/01/2002	$\Diamond$	

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application No.	An	plicant(s)				
	Office Action Summary	10/003,132		X ET AL.				
	· · · · · · · · · · · · · · · · · · ·	Examiner		Unit				
	The MAILING DATE of this communication a	Christopher Nicho		· ·				
Period for Reply								
THE - Exte after - If th - If NO - Failt - Any	MAILING DATE OF THIS COMMUNICATION PERIOD FOR REP MAILING DATE OF THIS COMMUNICATION ensions of time may be available under the provisions of 37 CFR or SIX (6) MONTHS from the mailing date of this communication. The period for reply specified above is less than thirty (30) days, a result of the provision of the provision of the period for reply is specified above, the maximum statutory period to reply within the set or extended period for reply will, by state reply received by the Office later than three months after the mailed patent term adjustment. See 37 CFR 1.704(b).	1.  1.136(a). In no event, however  eply within the statutory minim  d will apply and will expire SI.  ute, cause the application to be	er, may a reply be timely fil um of thirty (30) days will K (6) MONTHS from the m ecome ABANDONED (35	ed  pe considered timely.  ailing date of this communication.  U.S.C. § 133).				
1) 🖂	Responsive to communication(s) filed on 13	3 December 2001						
2a)[		This action is non-fina	اد					
3)	Since this application is in condition for allow			cution as to the morits is				
, —	closed in accordance with the practice unde							
4)⊠	Claim(s) 1-19 is/are pending in the application	on.						
	4a) Of the above claim(s) is/are withdrawn from consideration.							
5)[	Claim(s) is/are allowed.							
6)[								
7)	Claim(s) is/are objected to.							
8)⊠	8) Claim(s) <u>1-19</u> are subject to restriction and/or election requirement.							
Applicat 	ion Papers							
·	The specification is objected to by the Examir							
10)	The drawing(s) filed on is/are: a)☐ acc	•	•					
445	Applicant may not request that any objection to t							
11)	The proposed drawing correction filed on		. —	by the Examiner.				
If approved, corrected drawings are required in reply to this Office action.								
	The oath or declaration is objected to by the E	examiner.						
	ander 35 U.S.C. §§ 119 and 120							
	13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).							
a)	a) All b) Some * c) None of:							
	1. Certified copies of the priority documents have been received.							
	2. Certified copies of the priority documents have been received in Application No							
* 5	<ul> <li>3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).</li> <li>* See the attached detailed Office action for a list of the certified copies not received.</li> </ul>							
14) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).								
a) The translation of the foreign language provisional application has been received.  15) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.								
ر ریارہ Attachmen		sao priority unuer 33	0.0.0. 33 120 and	/Ot 12 1.				
1)  Notic 2)  Notic	e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTO-948) nation Disclosure Statement(s) (PTO-1449) Paper No(s)	5) 🔲 N		0-413) Paper No(s) Application (PTO-152)				

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## **DETAILED ACTION**

## Election/Restrictions

- 1. Restriction to one of the following inventions is required under 35 U.S.C. 121:
  - I. Claims 1-8 and 18, drawn to an isolated polypeptide, classified in class 530, subclass 350, for example.
  - II. Claims 9-17, drawn to method for producing a polypeptide comprising an isolated polynucleotide, expression vectors, and host cells comprising the same, classified in class 435, subclass 70.1, for example.
  - III. Claim 19, drawn to an antibody, classified in class 530, subclass 387.1, for example.
- 2. The inventions are distinct, each from the other because of the following reasons:
- 3. Although there are no provisions under the section for "Relationship of Inventions" in M.P.E.P. § 806.05 for inventive groups that are directed to different products, restriction is deemed to be proper because these products constitute patentably distinct inventions for the following reasons. Inventions I and III are directed to products that are distinct both physically and functionally, are not required one for the other, and are therefore patentably distinct.

  Although the polypeptide of Invention I can be obtained using the antibody of Invention III it can be isolated using materially different methods, such as chemical synthesis or isolation from natural sources. Although the antibody of Invention III can be used to obtain the polypeptide of Invention I it can also be used in materially different methods, such as in various diagnostic (e.g., as a probe in immunoassays or immunochromatography), or therapeutic methods.

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4. Inventions II and I are related as process of making and product made. The inventions are distinct if either or both of the following can be shown: (1) that the process as claimed can be used to make other and materially different product or (2) that the product as claimed can be made by another and materially different process (MPEP § 806.05(f)). In the instant case the polypeptides of inventions I can be made using materially different methods such as chemical synthesis or isolation from natural sources.

- 5. Inventions III and II are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the different inventions of Inventions III and II are unrelated product and method, wherein each is not required, one for another. For example, the claimed method of Invention II does not recite the use or production of the antibody of Invention III.
- 6. FURTHERMORE, restriction to one of the following inventions is required under 35 U.S.C. 121:
  - A. Claims 1-19, each in part, as the inventions pertain to SEQ ID NO: 2.
  - B. Claims 1-19, each in part, as the inventions pertain to SEQ ID NO: 4.
- 7. The inventions are distinct, each from the other because of the following reasons:
- 8. Although there are no provisions under the section for "Relationship of Inventions" in M.P.E.P. § 806.05 for inventive Inventions that are directed to <u>different</u> products, restriction is deemed to be proper because these products appear to constitute patentably distinct inventions

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for the following reasons: Inventions A and B are directed to sequences that are distinct both

physically and functionally, and are not required one for the other. Invention A requires search

and consideration of SEQ ID NO: 2, which is not required by the other Invention. Invention B

requires search and consideration of SEQ ID NO: 4, which is not required by the other Invention.

Each sequence requires a separate search of the literature and sequence databases. A search and

examination of an Invention as it pertains to all sequences would therefore present the examiner

with an undue search burden.

9. Applicant is advised that this is not a requirement to elect a species. Rather, this is a

second restriction requirement superimposed upon the requirement to elect one group from I-III.

In order to be fully responsive, Applicant must elect one group from I-III and one group from A-

B.

10. This application contains claims directed to the following patentably distinct species of

the claimed invention:

- a. Maltose binding protein
- b. An immunoglobulin constant region
- c. A polyhistidine tag
- d. A peptide as shown in SEQ ID NO: 7
- e. A peptide linker consisting of up to 25 amino acid residues

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 3 and 13 are generic.

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If applicant selects Invention I or II, one species from the second polypeptide group must be chosen to be fully responsive.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

- 11. Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).
- 12. Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter, separate search

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requirements, and/or different classification, restriction for examination purposes as indicated is proper.

13. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

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Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Christopher Nichols, Ph.D. whose telephone number is 703-305-

3955. The examiner can normally be reached on Monday through Friday, 8:30AM to 5:00PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's

supervisor, Gary Kunz, Ph.D. can be reached on 703-308-4623. The fax phone numbers for the

organization where this application or proceeding is assigned are 703-872-9306 for regular

communications and 703-872-9307 for After Final communications. The fax phone numbers for

the customer service center is 703-872-9305.

Any inquiry of a general nature or relating to the status of this application or proceeding

should be directed to the receptionist whose telephone number is 703-308-0196.

CJN

October 31<sup>st</sup>, 2002

Elyaber C. Kemmeres

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PRIMARY EXAMINER